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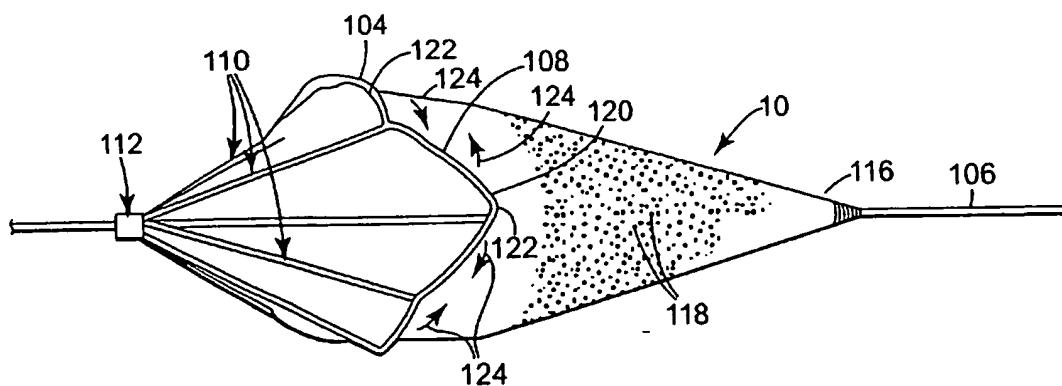
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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: MEDICAL DEVICES COMPRISING NANOMATERIALS AND THERAPEUTIC METHODS UTILIZING THE SAME



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(57) Abstract: The present invention provides medical devices comprising nanomaterials. By utilizing nanomaterials in the production thereof, the inventive medical devices can be provided with various advantageous properties and/or capabilities. Methods of producing the inventive medical devices are also provided. Inasmuch as the inventive devices are expected to provide certain advantages in their use, there is also provide a method of medical care, including methods of treatment or diagnosis, wherein the inventive devices are brought into therapeutic contact with a body to be treated or diagnosed thereby.

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61F2/02

A61F2/06

A61F2/01

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61F A61L A63B A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 6 228 471 B1 (PERSOONE PETER ET AL) 8 May 2001 (2001-05-08) column 4, line 29 -column 7, line 16 ---	1-4,6, 12,13, 30-32
X	WO 00 57932 A (BRIGHAM & WOMENS HOSPITAL) 5 October 2000 (2000-10-05)	1,2,30
A	page 4, line 17 -page 17, line 8 ---	12
A	GB 2 285 401 A (SUMITOMO RUBBER IND) 12 July 1995 (1995-07-12) page 2, line 17 -page 4, line 18 ---	8-10
P,X	WO 01 87193 A (SUPRONOWICZ PETER ;BIZIOS RENA (US); AJAYAN PULICKEL (US); SIEGEL) 22 November 2001 (2001-11-22) paragraph '0009! - paragraph '0019! ---	1-4,6, 12-18, 30-32
	-/-	

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

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20 March 2003

28/03/2003

Name and mailing address of the ISA

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X, P	US 2002/014182 A1 (VIGLIOTTI ANTHONY ET AL) 7 February 2002 (2002-02-07) paragraph '0036! – paragraph '0037! -----	1-4, 6, 12-18, 30, 31

INTERNATIONAL SEARCH REPORT

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: 19-29 because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery
2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-11,30-37

A medical device comprising at least one nanomaterial

2. Claims: 12-18

A medical care kit, comprising at least one medical device and at least one nanomaterial provided in packaging

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
US 6228471	B1	08-05-2001	EP WO AT DE DE EP ES JP	0856592 A1 9833948 A1 211510 T 69803365 D1 69803365 T2 0963455 A1 2171010 T3 2001509841 T	05-08-1998 06-08-1998 15-01-2002 28-02-2002 31-10-2002 15-12-1999 16-08-2002 24-07-2001
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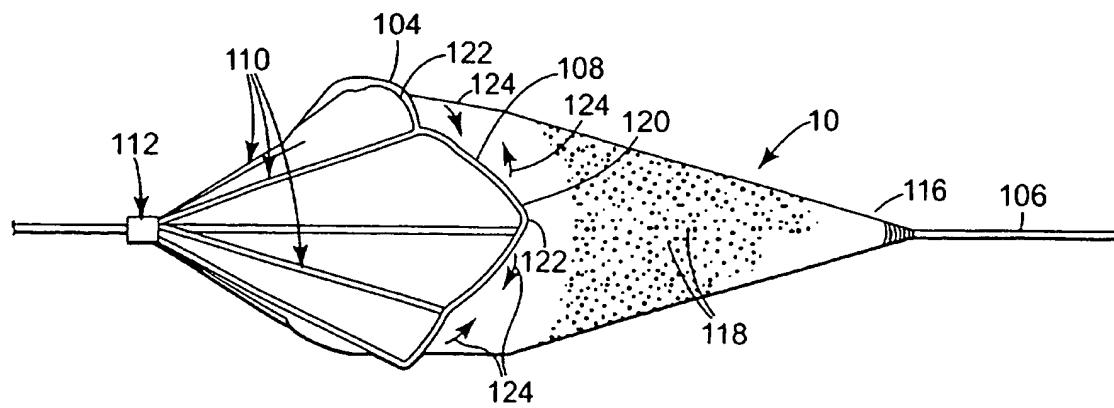
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(72) Inventor: WEBER, Jan; 18112 89th Place North, Maple Grove, MN 55311 (US).

(74) Agents: JORDAHL, Kimberly, S. et al.; Kagan Binder, PLLC, Suite 200 Maple Island Building, 221 Main Street North, Stillwater, MN 55082 (US).

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(54) Title: MEDICAL DEVICES COMPRISING NANOMATERIALS AND THERAPEUTIC METHODS UTILIZING THE SAME



WO 03/026532 A2

(57) Abstract: The present invention provides medical devices comprising nanomaterials. By utilizing nanomaterials in the production thereof, the inventive medical devices can be provided with various advantageous properties and/or capabilities. Methods of producing the inventive medical devices are also provided. Inasmuch as the inventive devices are expected to provide certain advantages in their use, there is also provide a method of medical care, including methods of treatment or diagnosis, wherein the inventive devices are brought into therapeutic contact with a body to be treated or diagnosed thereby.

5 **MEDICAL DEVICES COMPRISING NANOMATERIALS AND
THERAPEUTIC METHODS UTILIZING THE SAME**

The present invention relates to medical devices including one or more components comprising one or more nanoparticles and/or nanocomposite materials. By utilizing these nanomaterials in the manufacture of the inventive medical devices, certain properties of the nanoparticles and/or nanocomposites may be exploited in ways particularly advantageous in the medical device industry.

The medical device industry is but one example of an industry where the products or devices produced and used therein requires the products to exhibit a diverse array of properties and/or capabilities. Transluminal medical devices are one example. Such devices are typically introduced into the vasculature of a patient at a point remote from the treatment site, a procedure that can be uncomfortable for the patient. In order to perform acceptably, and to minimize the trauma to the patient, transluminal devices typically exhibit diverse, and at times divergent, performance characteristics and perform a wide range of functions.

For example, many such devices desirably exhibit good maneuverability so as to be manipulated to and/or inserted at a location requiring treatment, but yet sufficiently strong in the longitudinal direction so as not to buckle or kink when being so manipulated. In fact, many medical devices require a combination of these, and other, properties such as strength, thermal stability, structural stability, flexibility, opacity, radio-opacity, storage stability, lubricity, stability to sterilization treatment, etc., in order to be effective for their intended purpose. Often medical devices are further desirably biodegradable, capable of delivering therapeutic agents, etc.

30 Material selection is thus very important to the therapeutic efficacy of many medical devices since the properties of the materials used often dictates the properties and/or capabilities of the overall device. However, the range of properties available from one, or even a combination of, material(s) is often not as broad as

- 2 -

would be desired to provide a corresponding breadth of properties or capabilities in medical device applications. As a result, many medical devices need to be manufactured from a combination of materials, processed in a specific manner, or subjected to other treatments, in order to exhibit the desired and/or required

5 characteristics.

Thus, there is a continuing need in the medical device industry to develop or discover additional materials that exhibit the range of properties or capabilities required or desired for a medical device.

10 The present invention provides medical devices comprising nanoparticles and/or nanocomposite materials. According to the invention, the utilization of such nanomaterials in the manufacture of medical devices can provide the inventive devices with many, or all, of the diverse properties and/or capabilities often desirable in the same. That is, inasmuch as such devices often desirably exhibit a

15 vast number of often times divergent, properties, and/or desirably perform a variety of functions, it can be difficult to manufacture such devices without utilizing an extensive number of materials and processing techniques. By employing the present invention however, medical devices can be produced with a desired array of properties and/or capabilities using a lesser amount of materials and/or processing

20 techniques, or medical devices can be produced wherein one or more of the properties and/or capabilities are enhanced.

As a result, the present invention provides a medical device comprising at least one nanomaterial. The nanomaterial, in turn, may comprise at least one plurality of nanoparticles and/or at least one nanocomposite. The nanomaterial(s) may be present in one or more components of the device, or may be utilized to produce the device in total. Further, the nanomaterial(s) may be present as a tie layer, a coating or any other layer of the component or overall device. In certain advantageous embodiments, the nanoparticles may comprise a therapeutic agent.

Exemplary medical devices to which the invention is particularly directed

30 include balloons, catheters, shafting, filters and stent delivery systems such as disclosed in U.S. Patent Nos. 5,843,032; 5,156,594; 5,538,510; 4,762,129;

- 3 -

5,195,969; 5,797,877; 5,836,926; 5,534,007; 5,040,548; 5,350,395; 5,451,233;
5,749,888; 5,980,486; and 6,129,708.

Also provided is a method of making the inventive medical devices wherein the method comprises selecting a nanoparticle and/or nanocomposite material and 5 preparing at least a component of the medical device from the nanomaterial. The method may optionally include the steps of identifying the desired components of the nanoparticle and/or nanocomposite and preparing the nanomaterial according to any known technique.

The inventive medical devices can have enhanced properties and/or 10 capabilities relative to or properties and/or capabilities absent from, a corresponding medical device not comprising a nanomaterial. As a result, the inventive medical devices can provide certain advantages in their use. In this regard, the present invention also provides a method of treatment or diagnosis comprising bringing a medical device into therapeutic contact with a body to be treated or diagnosed, 15 wherein the medical device comprises at least one nanomaterial.

The accompanying drawings illustrate several aspects of the invention and together with description of the embodiment reserve to explain the principles of the invention. A brief description of the drawings is as follows:

20 Figure 1 is a longitudinal view, taken partially in cross-section, of a medical device in accordance with the present invention wherein the medical device is shown in a closed profile for insertion into a bodily lumen;

Figure 2 is a longitudinal view, taken partially in cross-section, of a medical device in accordance with the present invention wherein the medical device is 25 shown in an open, radially expanded profile;

Figure 3 is a longitudinal cross-sectional view of the distal end of a medical device in accordance with the present invention;

Figure 4 is a transverse cross-sectional view of the device shown in Figure 3, taken at line 4-4.

30

The embodiments of the present invention described below are not intended to be exhaustive or to limit the invention to the particular embodiments disclosed in

the following detailed description. Rather, the embodiments are described so that others skilled in the art understand the principles and practices of the present invention.

The nature of the production and therapeutic use of many medical devices
5 can require that the devices exhibit a broad and diverse array of properties and/or be capable of performing many functions. In order to achieve such a combination of desired properties and/or capabilities, more than one type of material is often employed in the construction of medical devices. For example, reinforcing particles can be applied or added to a substrate or matrix material to form a composite
10 material having a desired property, such as strength or an adhesion quality, otherwise lacking at the surface, or within the body of a matrix material.

Conventionally, the particles used in such composites are comprised of glass fibers, aggregates of amorphous or graphitic carbon, metal flakes, etc, and are at least about 1 micrometer in diameter in their largest dimension, or larger. While such
15 composite materials are useful in many medical device applications, the allowed tolerances in the manufacture of many other medical devices may not accommodate conventional, large size, filler particles.

Recently, a new class of particles has been described having at least one dimension less than about 1 micrometer. It has now been discovered that these
20 much smaller particles are particularly advantageously employed in medical device applications. The present invention thus provides medical devices comprising one or more of these nanomaterials.

As used herein, the term 'nanoparticle' is meant to indicate a particle having a greatest dimension of less than one micrometer. The nanoparticles may be
25 comprised of one or more materials, may be formed by using any processing technique, and may have any geometry, i.e., the nanoparticles can be generally spherical, octagonal or hexagonal, or they may be in the form of nanotubes, nanobelts, nanofibers, nanowires, and may be solid, hollow or porous, depending upon the desired application. Typically, the nanoparticles suitable for use in the
30 inventive medical devices, either alone or as a component of a nanocomposite material, have at least one dimension less than about 1000nm in size. In some embodiments, the nanoparticles can advantageously have one dimension of less than

about 750nm, typically less than about 500nm, for example, from about 1nm to about 100nm.

The term 'nanocomposite' generally refers to a composite material comprising at least one plurality of nanoparticles operatively disposed relative to a matrix material. The plurality of nanoparticles may be dispersed throughout the matrix material, either uniformly or otherwise, or may be substantially aggregated relative to a surface of the matrix material. Advantageously, the load or density of nanoparticles so disposed relative to the matrix material may be caused to vary across one or more dimensions of the matrix material as incorporated within the inventive medical devices, so that the property or capability enhanced or imparted by virtue of the use of the nanocomposite material within the inventive device can correspondingly vary. Finally, the term 'nanomaterial' is intended to indicate either or both nanoparticles and nanocomposites. That is, a medical device in accordance with the present invention comprising at least one nanomaterial may comprise only a plurality of nanoparticles, only a nanocomposite, or both.

For example, a medical device in accordance with the present invention wherein the nanomaterial comprises at least one plurality of nanoparticles would include a plurality of nanoparticles further comprising a therapeutic agent and a magnetic material and/or a heat sensitive thermoplastic material provided in combination with a device capable of attracting and/or adhering the nanoparticles thereto. Advantageously, this embodiment of the inventive medical device allows for the delivery of a therapeutic agent systemically, that can yet be delivered to, and substantially remain at, a desired treatment site at any desired time.

For example, such a plurality of magnetized nanoparticles could be provided with a magnetized device, such as a stent, or graft, that is itself magnetized, coated or wrapped with a magnetic material, such as a magnetized wire, or operably disposed relative to one or more magnets or other magnetized materials. Such a magnetized stent or graft could be delivered and or inserted at a desired treatment site and prior to, during, or at any time after, delivery of the stent or insertion of the graft, the magnetized nanoparticles could be caused to bind to the stent or graft via magnetic forces. For example, the magnetized nanoparticles could be injected as a

solution local to the stent or graft, could be injected systemically, or could be delivered via a perfusion balloon catheter.

Advantageously, such a delivery of magnetized nanoparticles could be caused to occur at any time during the resident life of the stent or graft. Further 5 advantages of this embodiment of the invention include the ability to systemically and yet directedly deliver any amount or number of therapeutic agents, as well as the ability to deliver such agents to a treatment site comprising a device that may not otherwise be incorporated into the device, e.g., therapeutic agents that are sterilization sensitive, such as DNA, proteins, viruses, and the like. Finally, just as 10 this embodiment of the invention allows for the systemic delivery of therapeutic agents at any time, it similarly is envisaged that this embodiment of the invention could allow for the removal of therapeutic agent from the treatment site. That is, a collection device capable of generating a stronger magnetic field than that of the device could be removably delivered to the treatment site to attract and adhere the 15 magnetized nanoparticles, and thus, any remaining therapeutic agent.

Further, in this embodiment of the invention, the nanoparticles may further comprise a heat sensitive thermoplastic material, either in place of, or in combination with the magnetic material. The desired device could then be operatively disposed relative to an energy source, so that the nanoparticles may be 20 systemically delivered, and once carried to the desired treatment site the energy source may be activated thereby heating the device and/or the nanoparticles. As a result, the heat sensitive material included in the nanoparticles can be caused to soften and adhere to at least a portion of a surface of the device.

Yet another example wherein an inventive medical device may comprise a 25 plurality of nanoparticles, not necessarily incorporated within or disposed relative to a matrix material, would include those embodiments of the invention wherein the nanoparticles are provided as a 'tie-layer' between two layers of the inventive device. One particular example of such an embodiment would include the utilization of carbon or ceramic nanotubes or fibers as a tie layer between two layers 30 that are desirably heat welded, coextruded, spun, sprayed or otherwise caused to be bonded together. In particular, a layer of such nanoparticles may be deposited on a first layer, either randomly, as may occur by dipping a first layer into a solution of

such nanoparticles, or in a relatively patterned or uniform fashion, such as a braid, as may be formed by spinning or spraying the nanoparticles on the first layer or surface. The second layer would then be deposited as desired, and via the method of application of the second layer or the subsequent application of heat, as by e.g., heat 5 welding, portions of the nanoparticles would become at least partially intertwined in either or both of the first and second layers of material, the resulting bond between the first and second layers being strengthened thereby. Clearly, this embodiment may also be encompassed within that embodiment of the invention wherein the medical devices comprise at least one nanocomposite materials, inasmuch as the 10 uniform dispersion, or in fact any dispersion, of nanoparticles throughout a matrix material is not required to fall within the definition of the term 'nanocomposite' as that term is used herein.

The nanomaterial to be used in the present medical devices is not particularly restricted. Rather, any nanomaterial that can be prepared or engineered to provide or 15 enhance at least one property desired in the desired medical device, or to provide a medical device with a desired capability, can be used. The selection of the particular nanoparticles, or the nanoparticles and matrix material(s), for use in the nanomaterial(s) will depend on the intended use of the medical device into which the nanomaterial will be incorporated and the desired properties of a device to be used 20 in that manner.

Additionally, the manner in which the nanocomposite is incorporated into the overall device can further be engineered and optimized in order to achieve or enhance desired properties and/or capabilities in the inventive medical device. For example, in a multilayered device, one layer of the device could be a nanomaterial, 25 with the remaining layers being comprised of traditionally filled materials, non-composite materials, or a combination of these. Further, the number and organization of the layers can be chosen to optimize and/or to provide properties desired in the device.

Generally speaking then, a matrix material according to the invention may 30 be any material suitable, or later determined to be suitable, for use in such a medical device. The matrix material may be any material that is historically or currently utilized, or contemplated for future use, in a corresponding medical device not

comprising a nanocomposite component. The matrix material may be comprised of organic, inorganic or hybrid organic/inorganic materials. Additionally, the matrix material may be a single material or a combination of materials, e.g., the matrix material may be a metal alloy, copolymer or polymer blend.

5 Exemplary matrix materials include, for example, polymers, such as thermoplastics and thermosets. Examples thermoplastics suitable for use as a matrix material include, for example polyolefins, polyamides, polyesters, polyethers, polyurethanes, polyureas, polyvinyls, polyacrylics, fluoropolymers, copolymers and block copolymers thereof, and mixtures thereof. Representative examples of
10 thermosets that may be utilized as a matrix material include elastomers such as EPDM, epichlorohydrin, nitrile butadiene elastomers, silicones, etc. Conventional thermosets such as epoxies, isocyanates, etc., can also be used. Biocompatible thermosets may also be used and these include, for example, biodegradable polycaprolactone, poly (dimethylsiloxane) containing polyurethanes and ureas, and
15 polysiloxanes.

 Analogously, nanoparticles according to the invention may be comprised of any material, or combination of materials, suitable, or later determined to be suitable, for use in a medical device, so long as the greatest dimension of the nanoparticles measures 1000nm or less, typically 750nm or less, more typically
20 500nm or less, for example, from about 1nm to about 100nm. The nanoparticles may comprise any material that has been, is, or is contemplated for use, as a conventionally sized filler material in a medical device. Desirably, the nanoparticles comprise a material capable of at least minimally altering the physical, mechanical, chemical, or other, properties of a matrix material when incorporated therein, or of
25 enhancing or providing the inventive medical device with a desired capability.

 As such, the nanoparticles may be comprised of any one or combination of organic, inorganic, or hybrid organic/inorganic materials. Additionally, the nanoparticles may be a single material or a combination of materials. For example, the nanoparticles may comprise a metal alloy, copolymer or polymer blend, or may
30 comprise at least a biocompatible and biodegradable material and a therapeutic agent. This latter embodiment may be advantageously employed in those

applications where the systemic delivery of therapeutic agents to a treatment site is deserved.

Examples of materials suitable for use in the nanoparticles include, among others, synthetic or natural smectic phyllosilicates including clays and micas (that 5 may optionally be intercalated, exfoliated and/or otherwise chemically modified) such as montmorillonite (mmt), hectorites, hydrotalcites, vermiculite, and laponite; monomeric silicates such as polyhedral oligomeric silsequioxanes (POSS) including various functionalized POSS and polymerized POSS; carbon and ceramic nano-tubes, nano-wires and nano-fibers, including carbon nanofibers and nanotubes of 10 any geometry prepared by electrospinning a carbonizable material, such as polyacrylonitrile; single and multi-walled fullerene nanotubes, silica nanogels, and alumina nano-fibers, as well as metal and metal oxide powders including aluminum oxide (AlO_3), titanium oxide (TiO_2), tungsten oxide, zirconium oxide, gold (Au), silver (Ag), platinum (Pt) and magnetic or paramagnetic materials such as 15 neodinium iron boron or super paramagnetic ferrite oxide (Fe_3O_4) or super paramagnetic maghemite (Fe_2O_3); organic materials including temperature sensitive polymers, such as polyvinylpyrrolidone and n-isopropylacrylamide copolymers or blends, and poloxamer, biodegradable polymers such as poly(lactic) acid, polysaccharide, polyalkylcyanoacrylate, which biodegradable polymers may also be 20 magnetized; and further including polytetrafluoroethylene, and dendrimers or dendrimer metal complexes.

The amount of the nanoparticles, or combinations of nanoparticles comprised of different materials, to be incorporated into the matrix or inventive medical device can vary depending on the desired properties exhibited by a particular medical 25 device or medical device component. Generally speaking, enough of the nanoparticles should be included so that desired property(ies) and/or capability(ies) are at least minimally exhibited by the nanocomposite and/or the medical device. On the other hand, not so much of the nanoparticles should be included so as to have a detrimental effect on the properties of the nanocomposite and/or medical device.

30 With this in mind, and while the particular range may vary depending on the nanoparticles and matrix material being utilized, or further on the desired properties or capabilities of the inventive medical device, it is believed that a useful range of

nanoparticles for inclusion in the inventive medical devices is from about 0.005% weight to about 99% weight of the total weight of the nanocomposite or overall medical device, as the case may be. In many embodiments, nanoparticles may be incorporated in an amount of from about .01% up to about 40% or 50% by weight 5 and as a percentage of the total weight of the nanocomposite and/or medical device, as the case may be. In a typical embodiment, the nanoparticles can be incorporated in an amount of from about 0.1% to about 20% by weight of the nanocomposite or medical device, for example, from about 1% to about 10% by weight of the nanocomposite or medical device.

10 The present invention further contemplates that there may be applications in which it will be desirable to have a combination of more than one plurality of nanoparticles, so that each different plurality may be comprised of a different material or combination of materials. In this manner, a further enhancement of a single desired property, or an addition of a new property or capability may be seen 15 in the inventive medical device.

For example, it may be advantageous to prepare a nanocomposite comprising a polymeric matrix material and a first plurality nanoparticles that exhibits radio-opacity. The nanocomposite, or inventive device itself, may desirably further comprise a second plurality of nanoparticles that comprises a strength enhancing 20 material. A device according to this embodiment of the invention would not only be visible via conventional imaging techniques, but further, could exhibit enhanced strength relative to a corresponding devices not comprising the strength enhancing nanoparticles.

As but one particular example of this embodiment of the invention, a stent 25 could be provided that is coated with a nanocomposite prepared with an appropriate matrix material and a first plurality of magnetic nanoparticles. A second plurality of nanoparticles comprising a magnetized biodegradable, biocompatible material, such as magnetized polylactic acid, and a therapeutic agent could also be provided. As described above, the stent could then be implanted at the desired treatment site 30 and the second plurality of particles could be delivered systemically either concurrent with, or subsequent to, the delivery of the stent. The magnetic materials of the stent and the second plurality of nanoparticles could attract each other and

adhere at least a portion of the second plurality of nanoparticles to the stent. As also described above, it is not necessary that the stent comprise magnetic nanoparticles, or any nanomaterial at all, rather small magnets could be operatively provided in connection with the stent.

5 This embodiment thus not only provides for the targeted and perhaps even controlled delivery of therapeutic agents to a desired treatment site, but further, provides the opportunity to systemically deliver the therapeutic agent at any desired time. That is, whereas prior art medical devices achieve drug delivery by incorporating the drug in a timed release coating on the device, thereby limiting the 10 time within which the drug is delivered to the time that the device is utilized or implanted and for a short time period thereafter, this embodiment of the present invention would allow for the delivery of therapeutic agents to a desired treatment site long after the implantation of the inventive medical device.

15 In those embodiments of the present invention wherein the nanoparticles desirably comprise a therapeutic agent, any therapeutic agent may be incorporated relative thereto. The therapeutic agent can be genetic or non-genetic or may comprise cells or cellular matter. Examples of non-genetic therapeutic agents include, but are not limited to, antithrombogenic agents such as heparin and its derivatives, urokinase, and dextropheylalanine proline arginine chloromethylketone (Ppack); anti-proliferative agents such as enoxaprin, andiopeptin, or monoclonal antibodies capable of blocking smooth muscle cell proliferation, hirudin, and acetylsalicylic acid; anti-inflammatory agents such as desamethasone, prednisolone, corticosterone, budesonide, estrogen, sulfasalazine, and mesalamine; antineoplastic/antiproliferative/anti-miotic agents such as paclitaxel, 5-fluorouracil, 20 cisplatin, vinblastine, vincristine, epothilones, endostatin, angiostatine and thymidine kinase inhibitors; anesthetic agents such as lidocaine, bupivacaine, and ropivacaine; anti-coagulants such as D-Phe-Pro-Arg chloromethyl keton, an RGD peptide-containing compound, heparin, antithrombin compounds, platelet receptor antagonists, anti-thrombin antibodies, anti-platelet receptor antibodies, aspirin, 25 prostaglandin inhibitors, platelet inhibitors and tick antiplatelet peptides; vascular cell growth promoters such as growth factor inhibitors, growth factor receptor antagonists, transcriptional activators, and translational promoters; vascular cell 30

growth inhibitors such as growth factor inhibitors, growth factor receptor antagonists, transcriptional repressors, translational repressors, replication inhibitors, inhibitory antibodies, antibodies directed against growth factors, bifunctional molecules consisting of a growth factor and a cytotoxin, bifunctional molecules

5 consisting of an antibody and a cytotoxin; cholesterol-lowering agents; vasodilating agents; and agents which interfere with endogenous vasoactive mechanisms.

Examples of genetic materials that can be used as therapeutic agents include, but are not limited to, anti-sense DNA or RNA; and DNA coding for anti-sense RNA, tRNA or rRNA, e.g., to replace defective or deficient endogenous molecules,

10 angiogenic factors including growth factors such as acidic and basic fibroblast growth factors, vascular endothelial growth factor, epidermal growth factor, transforming growth factor α and β , platelet-derived endothelial growth factor, platelet derived growth factor, tumor necrosis factor α , hepatocyte growth factor and insulin like growth factor, cell cycle inhibitors including CD inhibitors, agents

15 useful for interfering with cell proliferation such as thymidine kinase, and the family of bone morphogenic proteins (BMP), or any other molecule capable of inducing an upstream or downstream effect of a BMP, including BMP-2, BMP-3, BMP-4, BMP-5, BMP-6 (Vrg-1), BMP-7 (OP-1), BMP-8, BMP-9, BMP-10, BMP-11, BMP-12, BMP-13, BMP-14, BMP-15 and BMP-16.

20 The properties and/or capabilities of the nanocomposites and/or inventive medical devices may be affected by compatibility of, and/or the level and/or kind of interaction that occurs between, the nanoparticles and the matrix material of the nanocomposite, or of the nanoparticles and the inventive medical device. For example, the nanoparticles and matrix material and/or the device may also interact 25 chemically, such as by the establishment of Van Der Waal's forces, covalent bonds or ionic bonds between the filler particles and the matrix material.

Or, the compatibility may be such that the nanoparticles and the matrix and/or medical device interact physically, such as by chain entanglement of the nanoparticles with the matrix material and/or the surface of the medical device via 30 the physical interaction of the device and the heat sensitive polymer. For example, in that embodiment of the invention wherein the nanoparticles comprise a heat sensitive polymer and a therapeutic agent, a source of heat external to a medical

- 13 -

device implanted within a host could be used to cause the heat sensitive polymer of the nanoparticle to soften and adhere to the medical device via the physical interaction of the device and the heat sensitive polymer. Said softening could also assist in the release of the therapeutic agent, and thus, such an inventive medical 5 device could be utilized to provide for targeted drug delivery. As another example, in that embodiment of the invention where the nanoparticles comprise a magnetic material, physical interaction could be caused to occur with the surface of a medical device via the provision of magnets operatively disposed relative to a surface of the medical device. If the magnetic material were further a biodegradable magnetic 10 material, and the nanoparticle further included a therapeutic agent, targeted, timed, and controlled drug delivery could be provided by the device.

Generally speaking, any such compatibility, and the resulting interaction, can act to enhance the dispersion of the nanoparticles within the matrix material of a nanocomposite and/or to enhance the disposition of the nanoparticles relative to the 15 medical device. If this is the case, and very generally speaking, the greater the compatibility, the greater the increased dispersion and/or enhanced disposition. Therefore, in applications where such greater dispersion or enhanced disposition would be desirable, the compatibility of, and resulting interaction between, the nanoparticles with the matrix material and/or the medical device can be encouraged 20 or facilitated.

The compatibility of the nanoparticles and the matrix material can be enhanced, for example, simply by selection of the materials for use as the matrix or in the nanoparticles. That is, interaction between the nanoparticles and the matrix may be facilitated or caused simply by selecting nanoparticles and matrix materials, 25 and/or other medical device material(s), with compatible functional groups. If such compatible functional groups are not present, they can be provided by 'functionalizing' the nanoparticles matrix material, or desired surface of the medical device to provide compatible functional groups that can then interact with each other.

30 Smectic phyllosilicates, monomeric silicates, and ceramics are just a few representative examples of materials suitable for use in the nanoparticles that may be advantageously functionalized to provide increased interaction between the

nanoparticles and the matrix material and/or medical device. For example, POSS monomers can be functionalized with, e.g., organic side chains to enhance compatibility with, e.g., polystyrene. Furthermore, clays, such as montmorillonite may be functionalized with alkylammonium so that the montmorillonite is capable 5 of interacting with a polyurethane, for example. Clays may further be functionalized with block or graft copolymers wherein one component of the copolymer is compatible with the clay and another component of the copolymer is compatible with the polymer matrix.

In addition to functionalizing any or all of the nanoparticles, the matrix 10 material, or the surface of the inventive device itself, the compatibility of, and interaction between, the nanoparticles, matrix material and/or medical device can be enhanced by incorporating one or more coupling or compatibilizing agents into the nanocomposite, or onto the surface of the inventive device itself. As used herein, the term(s) coupling/compatibilizing agents include any agent capable of enhancing 15 compatibility and/or promoting interaction between the nanoparticles and the matrix material, or between the nanoparticles and the inventive device.

Such agents can be organic or inorganic. The selection of these optional 20 agents will, of course, depend on the nanoparticles, matrix material, and/or the nature of the portion of the inventive device with which the nanoparticles desirably interact. Bearing this in mind, suitable organic coupling agents can be both low molecular weight molecules and polymers. Examples of low molecular weight 25 organic coupling/compatibilizing agents include, but are not limited to, amino acids and thiols. Examples of polymeric compatibilizers would include functionalized polymers, such as maleic anhydride containing polyolefins or maleimide-functionalized polyamides. Inorganic coupling agents would include, for example, alkoxides of silicon, aluminum, titanium, and zirconium, or magnetic powders such as ferrite oxide, to name a few.

Generally speaking, the amount of a coupling/compatibilizing agent used, if 30 used at all, will desirably be that amount which will at least marginally improve the compatibility of the nanoparticles and the matrix material, or the nanoparticles and the inventive medical device itself, so that at least a minimal enhancement of the dispersion of the nanoparticles within the matrix and/or the properties of the

nanocomposite, or an enhanced interaction between the nanoparticles and the inventive device, can be observed. Useful amounts of such agents are contemplated to be within the ranges of from about 0.1% to about 10% by weight of the nanocomposite; typically from about 0.5% to about 7.5%, more typically from about 5 1% to about 5% by weight of the nanocomposite.

In addition to material selection, functionalizing and compatibilizing as a means to promote interaction of the nanoparticles and the matrix material, said interaction may be enhanced, if desired, by utilizing compatibility enhancing methods of processing such as ultrasonic assisted extrusion and/or compounding.

10 By applying an ultrasonic vibration to the extruder die, for example, the friction shear forces can be reduced, and a melt comprising the matrix material and nanoparticles rendered more homogeneous. Additionally, the nanoparticles and/or matrix material can be dispersed in a solvent, e.g., dimethylformamide, dichloroethylene, N-methyl-2-pyrrolidone and the like. Once so dispersed, the 15 nanoparticle solution and the matrix material solution could be mixed and applied onto a mandrel via electro, or electrohydrodynamic spraying or spinning, or any other suitable method to produce a nanocomposite material or inventive medical device, exhibiting enhanced dispersion of the nanoparticles. Any other techniques of enhancing the dispersion of nanoparticles within a matrix can also be utilized, if 20 such an enhanced dispersion is desirable in the chosen application.

While it may be desirable in certain applications to increase the interaction between the nanoparticles and the matrix material, or between the nanoparticles and the device itself, extensive interaction between the nanoparticles themselves can be undesirable in certain applications. In particular, in applications where the 25 nanoparticles desirably form a layer with a substantially uniform thickness, or where an otherwise substantially uniform dispersion throughout a matrix material or relative to a medical device is desired, any substantial agglomeration of the nanoparticles can be suboptimal. In such applications then, it may be advantageous or desirable to include a dispersant in solution with the nanoparticles prior to their 30 dispersion within or application to, the matrix material and/or the inventive device.

In addition to the nanoparticles, the matrix material, and optionally, coupling/compatibilizing agents or dispersants, the nanocomposites and/or medical

devices according to the invention can comprise any other materials utilized in a corresponding medical device not comprising a nanomaterial. For example, pigments and/or whiteners, can be provided. Also, processing aids, such as plasticizers, surfactants and stabilizers can be included in the nanocomposites. Also, 5 conductive, magnetic and/or radiopaque agents could be included. Such agents, the amounts in which they are useful, as well as the benefits that they provide, are well known to those of ordinary skill in the art.

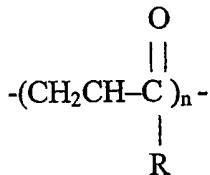
One example of a class of stabilizers that may find use in the inventive medical devices and methods is that commonly referred to as radiation oxidative 10 degradations, or "ROD" stabilizers. As the name suggests, these agents may assist a polymer within which they are incorporated to resist any degradation that may otherwise occur upon exposure of the polymer to sterilizing radiation. Additionally, however, such stabilizers may also be useful in assisting a polymer to resist any degradation that may otherwise occur during processing, such as during mixing 15 and/or heating that may be required in order to adequately disperse nanoparticles throughout a matrix material.

Such ROD stabilizers may be antioxidants, particularly radical or oxygen scavengers. Mercapto compounds, hindered phenols, phosphites, phosphonites and hindered amine antioxidants are among the most effective such stabilizers. Specific 20 examples of stabilizers are 2-mercaptobenzimidazole, triauryl phosphite, IONOX 330, 2-mercaptobenzothiazole, N,N-di(β -napthyl-*p*-phenylenediamine((DPPD), SANTONOX R, SANTOWHITE powder, phenothiazine, IONOL, 2,6-di-*t*-butylcresol, N-cyclohexyl-N'-phenyl-*p*-phenylenediamine, nickel 25 dibutyldithiocarbamate, IRGANOX 1010, β -(3,5-di-*t*-butyl-6-hydroxyphenyl) propionate, 1,2,2,6,6-pentamethyl-4-stearoyl piperidine, and 2,2,6,6, tetramethyl-4-nitropiperidine. Further examples include butylated reaction product of p-cresol and 30 dicyclopentadiene, substituted amine oligomers, N,N'-bis(2,2,6,6-tetramethyl-4-piperidinyl)-1,6-hexanediamine, 2,4-dichloro-6-(4-morpholinyl)-1,3,5-triazine, and N,N'-hexamethylene-bis[3-(3,5-di-*t*-butyl-4-hydroxyphenyl)propionamide]. Still further, transition metals or compounds thereof may function as ROD stabilizers, for instance iron, cobalt, nickel, ruthenium, rhodium, palladium, osmium, iridium,

platinum, copper, manganese and zinc metal and compounds, as described in WO 99/38914, US 5034252, and US 5021515.

The ROD stabilizer may also be an oxygen scavenging polymer, such as the polyketone polymers described in WO 96/18686 of the formula

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where R is H, an organic side chain or a silicon side chain, and n is a positive number greater than 2. Such polyketone ROD stabilizers are suitably employed in the thermoplastic composition in an amount of from 0.1 to about 10% by weight.

If their presence is desired, ROD stabilizers may be employed in the nanocomposites in any amount at least minimally effective in assisting in the resistance of the matrix material to degradation, i.e., in amounts of from about 0.01% to about 5%, suitably from about 0.1 to about 1%, for instance from 0.2% to 0.5%. The stabilizer can be compounded into the nanocomposite in the extrusion melt or in a separate compounding step prior thereto.

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Many nanocomposites and nanoparticles are commercially available. Additionally, many methods of producing such nanocomposites and/or nanoparticles are known, and any of these maybe utilized to produce nanocomposites and nanoparticles for incorporation into the inventive device. Many such methods are disclosed and described, for example, in "Nanocomposites 2001, Delivering New Value to Plastics", Executive Conference Management, June 25-27, 2001, Chicago, IL.

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Advantageously, and since the filler particles can have an impact on the properties exhibited by the nanocomposite by virtue of the dispersion of the filler particles within the matrix, the particular method utilized to prepare the nanocomposite can be selected to assist in the provision of medical device with the desired array of properties. That is, in certain medical device applications, it may be desirable to have the entirety of the medical device or medical device component exhibit the properties of the nanocomposite substantially uniformly throughout, or across the length of, the medical device. In such applications, it would be desirable

to substantially uniformly distribute the filler particles throughout the matrix of the nanocomposite. In other applications, it may be desirable to have the entirety of the medical device or medical device component exhibit the properties of the nanocomposite, but at varying degrees throughout the device or component. In 5 these applications, then, it would be desirable to vary the distribution of the filler particles throughout the matrix of the nanocomposite in a manner so that the desired varied properties are observed in the medical device or component.

For exemplary purposes only, then, processes for the production of such 10 nanocomposites include polymerization of the matrix material in the presence of the filler, melt compounding of the matrix with the filler, and in-situ formation of the filler, e.g., as would be provided by the adding a silane monomer to a block copolymer and then curing the silane to produce nanostructured silica filler particles relatively uniformly dispersed within the in the matrix of the copolymer, to name a few. If a coupling/compatibilizing agent is to be used, it may be pre-coated onto the 15 filler particles before compounding the filler particles with the matrix, or alternatively, the agents may be added during the nanocomposite formation process.

It has now been appreciated that these new nanomaterials can provide many 20 unique advantages in the production of medical devices, i.e., certain desirable properties of nanomaterials can be exploited in ways particularly advantageous in the medical device industry.

First, the use of nanomaterials in the manufacture of medical devices can provide a greater amount of control to the manufacturer over at least some of the properties of the resulting medical device. In fact, the use of a nanomaterial in the 25 inventive medical devices, either alone or in combination with one or more traditional materials, advantageously provides the opportunity to provide a variation in one or more properties within one or more dimensions of the inventive device than can be achieved in medical devices not comprising nanomaterials.

Further, a more subtle change in property and/or capability of a device can 30 be seen by making a small change in the amount of nanoparticles within the device than can be seen when utilizing conventionally sized filler particles. This change in amount or density can be caused to occur within the inventive device. That is, if a nanomaterial is being utilized, for example, to provide strength to a length of

medical device tubing or shafting, and it is desired that the strength of the tubing will decrease so that the tubing becomes more flexible at its proximal end, the load density of the nanoparticles within the length of tubing can be caused to change accordingly. Of course, such a load reduction can be caused to occur by reducing 5 the amount of nanoparticles within the nanocomposite along the length of the tubing, or by simply by altering the thickness of the tubing along its length.

Secondly, a lesser amount of filler material, relative to traditionally sized 10 filler particles, can produce a similar property or provide a similar capability. The use of a lesser amount of material can translate to less material, shipping and storage cost, a savings that can be an important advantage in many medical device applications. Even in applications where a greater concentration of filler particles is 15 desired, such highly filled nanocomposites are much easier to process than composites filled with the same concentration of conventionally sized particles.

Third, nanomaterials can, if desired, be comprised of materials traditionally 20 used in the manufacture of medical devices, so long as the nanoparticles have a greatest dimension of less than about 1 micrometer when processed from such materials. The inclusion of these nanomaterials into a medical device comprising other non-nanomaterials is thus not likely to cause a lack of cohesiveness in the overall device due to incompatibility between the nanomaterial and non-nanomaterials. As a result, if desired, a variation in one or more properties of a 25 multi-layered device could be provided by selecting and optimizing the number and organization of the layers, as well as the nanomaterials and non-nanomaterials to be used therein, to provide the properties or capabilities desired in the multilayer medical device.

Finally, the utilization of nanomaterials in the inventive medical devices 30 advantageously can be used to provide optically clear medical devices that are yet radiopaque. That is, since in many cases the size of the nanoparticle is smaller than the wavelength of visible light, it is possible to use nanocomposite materials to manufacture a device is transparent, and yet X-ray radiopaque. Other unique advantages that may be seen by the utilization of nanocomposites in the inventive medical devices include effects such as lowering the coefficient of friction, biocompatibility, and biodegradability, to name a few.

In light of the aforementioned newly identified and appreciated advantages the present invention thus provides medical devices including at least one component comprising at least one nanomaterial. The present invention contemplates that the nanomaterial may be utilized as one or more of a layers of a 5 component, a component of a device in total, a device in total, or as a coating deposited over any portion of any of these.

The applicability of the inventive concept is not particularly limited to any particular class or grouping of medical devices. Rather, the identified advantages that can be seen are believed to be widely applicable, in whole or in part, to any 10 medical device. It is believed, however, that the inventive concept will prove particularly advantageous when utilized in medical devices contemplated to be brought into therapeutic contact with a body, i.e., devices contemplated to be introduced into the body, either temporarily or permanently, for the purpose of effectuating a treatment or diagnosis thereof. Such devices find use in, e.g., urinary, 15 cardiovascular, musculoskeletal, gastrointestinal, or pulmonary applications.

Medical devices useful in urinary applications include, for example, catheters, shunts, stents, etc. Exemplary medical devices useful in cardiovascular applications include stents, angiography catheters, coronary or peripheral angioplasty catheters (including over the wire, single operator exchange or fixed 20 wire catheters), balloons, guide wires and guide catheters, real or artificial grafts, artificial vessels, artificial valves, filters, vascular closure systems, shunts, etc. Musculoskeletal medical devices include, for example, artificial ligaments, and prosthetics. One example of a medical device useful in a gastrointestinal application would be a shunt. Pulmonary medical devices include prosthetics, as one example.

25 Medical devices to which the inventive concept is believed to be particularly applicable include medical devices useful in any of the aforementioned applications, and that are introduced into the lumen of a patient. Such devices include, for example, catheters, embolic protection device, grafts, shunts, stents, balloons, etc., for use in urinary, cardiovascular, musculoskeletal, gastrointestinal or pulmonary 30 applications. Even more particularly, it is believed that the inventive concept will prove particularly advantageous when applied to medical devices intended to be introduced into a cardiovascular lumen for therapeutic purposes.

The present inventive medical device can, for example, be a catheter shaft, such as for an angiography system, angioplasty balloon, guide catheter, or stent delivery system. Such devices often include multiple lumens in a side-by-side or coaxial configuration. Coaxial configurations generally have inner and outer tubular components. According to the invention, the inner and/or outer tubular component of such a catheter can be formed from a nanomaterial. The inner and/or outer tubular components, in turn, can be comprised of a plurality of layers, in which case, one or more of the layers may comprise a nanomaterial. As mentioned hereinabove, and as is the case with any multi-layered medical device wherein at least one of the layers comprises a nanomaterial, the number and organization of the layers can be chosen and optimized to provide the properties or capabilities desired in the multilayer tubular component.

One particular example of a catheter shaft within the scope of the invention could comprise, for example, a nanocomposite further comprising carbon or ceramic nanofibers. Advantageously, such nanofibers could be utilized in applications wherein stainless steel braiding is currently utilized, thereby allowing for thinner walls than are possible with stainless steel braiding incorporated therein. Further, the utilization of ceramic or carbon nanofibers can result in a device that is detectable via MRI technology, whereas the presence of stainless steel braiding interferes with the image produced thereby. Finally, the increased surface area for contact between the nanofibers and the matrix material, as compared to the surface area for contact between a matrix material and stainless steel braiding, can result in a greater cohesion between the nanofibers and the matrix material.

Such shafting could be prepared by first spinning the nanofiber from the desired material, for example, by electrospinning or electrohydrodynamic spinning. Generally speaking, electrospinning uses electrical forces applied at the surface of a polymer solution or melt to create an electrically charged jet. When the jet dries or otherwise solidifies, an electrically charged fiber remains. This technique can be used to produce patterns or braids from the electrically charged polymer jet by directing the charged jet onto a rod or tube, while rotating either or both of the spray nozzle, or the rod or tube onto which the material is deposited. Any strength enhancing polymer may be so processed and utilized in the inventive shafting, and

polyacrylonitrile is but one example of a suitable polymer. Advantageously, and as is described in "Manufacturing, Characterization and Applications of Carbon Nanofibers", Yongkui Wen, Mat. Res. Soc. Symp. Proc., Vol. 702, the resulting polyacrylonitrile fibers can be carbonized to produce carbon fibers, which are from 5 20 to 100 times stronger than stainless steel fiber braiding. Once the nanofibers of the shafting have been so formed, a spray or dipcoat process can be used to provide the matrix material about the fiber network. Alternately, a nanocomposite comprising the nanofibers could be processed into the desired device utilizing normal extrusion, intermittent extrusion and/or by multi-layer extrusion.

10 Although described in connection with this particular exemplarily embodiment of the present invention, the utilization of nanofibers, as well as the utilization of the electrospinning or spraying process to form them, or any other nanoparticles, followed by dipcoating, spraying, etc., to provide a matrix material, or component of the inventive device, about the fibers, is clearly within the scope of 15 the present invention. For example, nanofibers, or a nanoparticle of any other geometry, form or shape, and comprising any material could be formed over a mold, and dipcoated or sprayed with, or welded to, a desirable matrix material, or component of an inventive device to produce any medical device to provide a medical device in accordance with the present invention.

20 A further example of a transluminal medical device that may benefit from application of the inventive concept would be a vessel graft. Such devices are known, and any of these, or vessel grafts developed in the future, may be improved by applying the inventive concept thereto. According to the invention, the graft may be artificial and may be prepared from, or coated with, a nanocomposite material, or 25 may be real tissue. In either case, the graft may have a plurality of nanoparticles operatively disposed relative to a surface of the graft.

For example, and as described briefly hereinabove, a graft may be provided having disposed operatively relative thereto a magnetic wire. A plurality of nanoparticles comprising, for example, a magnetized biodegradable polymer and a 30 therapeutic agent can be provided in conjunction with the graft. The graft can be implanted at a desired treatment site and the nanoparticles delivered systemically at any desired time to effectuate the delivery of the therapeutic agent to the treatment

site. More particularly, the nanoparticles, once delivered via intraluminal fluid flow to an area in proximity to the graft, will be attracted and adhere to the graft via the magnetic materials present therein. The degradation of the biodegradable polymer will cause the site specific release of the therapeutic agent over time.

5 Yet another example of a transluminal medical device useful in cardiovascular applications that can be advantageously enhanced via application of the inventive concept, is an embolic protection device, also known as a distal protection device, embolic filter, or simply as a filter. Many such devices are known, representative examples of which are described in commonly assigned U.S. 10 Patent Numbers 6,001,118 and 6,336,934.

Briefly then, and for purposes of illustration, embolic filters are utilized in transluminal cardiovascular applications wherein it is desired to collect floating debris, such as emboli. Generally, such filters have a collapsible body having a proximal inlet that allows blood, embolic material and/or other debris to enter the 15 filter and a distal outlet that allows the passage of blood therethrough, while yet retaining the embolic material and/or other debris within the filter body. Filters are desirably collapsible to a minimal profile for insertion into the luminal space from which emboli is desirably captured and removed, but yet, in order to be effective, desirably enlarge to approximate the inner diameter of the luminal space, i.e., so that 20 substantially all emboli at least enters the filter body. Thus, filters are typically comprised of a thin material to provide the desired low profile, and yet, this thin material must be sufficiently robust to withstand collapsing, insertion and expansion within a luminal space. Further, once so expanded, filters are desirably comprised of a material having sufficient strength to withstand the pressure exerted by blood 25 flowing and pulsing therethrough. Conventional filters, although acceptable, could yet be improved upon in these areas.

Application of the present inventive concept to embolic filters can provide such improvements. That is, by utilizing a nanomaterial in the production thereof, a thin filter can be produced that yet has sufficient elasticity, strength and robustness 30 to perform satisfactorily under its demanding conditions of use. As but one example of this particular embodiment of the invention, an embolic filter can be produced utilizing a nanocomposite comprising a matrix material comprising

polyurethane, such as Pellethane®, commercially available from the Dow Chemical Company, Midland, MI, and nanoparticles comprising functionalized clay. More specifically, a filter can be prepared by dissolving the polyurethane in a suitable solvent, such as tetrahydrofuran, adding the clay nanoparticles thereto, and mixing 5 the nanocomposite, e.g., in a high speed blender. The resulting nanocomposite can then advantageously be formed into an embolic filter by electrohydrodynamically spraying the nanocomposite onto a suitable mold or form. The resulting filter may advantageously comprise a substantially homogeneous dispersion of the clay nanoparticle due to the minimized droplet size distribution that can be seen utilizing 10 electrohydrodynamic spraying as opposed to conventional electrostatic spraying. Further, the resulting filter can be three times thinner and yet many times stronger than an embolic filter prepared from conventional materials.

Referring now to Figures 1 and 2, there is illustrated such an embolic filter embodying principles of the present invention. More particularly, Figures 1 and 2 15 illustrate embolic protection device 10 for collecting loosened debris in a body lumen, Figure 1 illustrating device 20 in a closed collapsed profile, for insertion into a bodily lumen, while Figure 2 illustrates device 10 in an open, radially expanded profile for collecting debris from a bodily lumen.

Device 10 includes a filter membrane 102 and a collapsible proximally 20 tapered frame 104. Frame 104 supports filter membrane 102 and is operably coupled to an elongated guidewire 106 or other support device. Frame 104 includes an orifice 108 and a plurality of longitudinally extending support ribs 110. In the expanded profile illustrated in Figure 2, orifice 108 is opened and support ribs 110 extend radially outwardly to support orifice 108. Collar 112 movably couples the 25 proximal ends of ribs 110 to guidewire 114. Orifice 108 is thus coupled to collar 112 through ribs 110 and is movable between the collapsed profile illustrated in Figure 1 and the opened deployed profile illustrated in Figure 2.

Filter membrane 102 is generally cone-shaped, having a proximal end 114 and distal end 116. Distal end 116 is preferably tapered so as to meet the diameter 30 of guidewire 106 and is preferably fixedly secured or formed to guidewire 106. Proximal end 114 has a larger diameter than distal end 116, preferably approximately equivalent to the diameter of orifice 108, and is coupled thereto.

According to the present invention, filter membrane 102 is formed from a nanocomposite material, such as the polyurethane/clay nanocomposite material discussed hereinabove, and has a plurality of small openings 118 provided therein. Openings 118 may be formed in the nanocomposite membrane by known laser 5 techniques and are desirably sized in order to allow blood to flow therethrough, while restricting the flow of debris or emboli, preferably retaining such debris or emboli within filter membrane 102.

Orifice 108 is generally formed of a pleated ring 120 having an expanded dimension to support filter membrane 102 in the opened deployed profile as 10 illustrated in Figure 1 and a collapse dimension to support the filter in the closed collapsed profile as illustrated in Figure 2. Pleated ring 120 is collapsed by closing folds 122 as illustrated by arrows 124 so that adjacent folds 122 are positioned in close proximity. In such a position, orifice 108 assumes a relatively small dimension to collapse filter membrane 102 for insertion into, and retrieval from, an 15 intraluminal space.

Referring now to Figures 3 and 4, there is illustrated an embodiment of a medical device embodying principles of the present invention. In particular, Figure 3 is a longitudinal cross-section view of the distal end of a balloon angioplasty catheter 30. In this embodiment, catheter 30 includes an inner tubular component 20 330 comprising an inner layer 332 and outer layer 334. A balloon 336 having a distal waist 338 is attached to inner tubular component 330. Balloon 336 also has a proximal waist 340 attached to outer tubular component 342. A guidewire 344 is shown within lumen 346 of inner tubular component 330. Figure 2 is a transverse cross-section view taken at line 2-2 of Figure 1.

According to the invention, it will be appreciated that inner tubular 25 component 330, including either or both of inner layer 332 and outer layer 334, outer tubular component 342, balloon 336, or guidewire 344, may be prepared from, or may comprise, a nanomaterial as disclosed herein. Furthermore, balloon 336 may have a stent (not shown) disposed operatively relative thereto, in order to provide 30 that catheter 30 may also function as a stent delivery system. In such an embodiment, the stent, as well as any other components that may desirably be included in such a stent delivery device, may comprise a nanomaterial.

In addition to the inventive devices themselves, the present invention also provides methods of making the inventive devices. Generally speaking, the method involves providing a nanomaterial and processing the nanomaterial to form at least a portion of at least a component of a medical device. The nanomaterial may be

5 obtained from any of a variety of commercial sources of such materials, or, it may be prepared from the desired components thereof by any method. The device or component thereof may then be manufactured or processed using the nanomaterial according to any suitable method, including at least those methods utilized to manufacture the corresponding non-filled medical device, or a corresponding

10 medical device comprising a composite material further comprising traditionally sized filler particles. There are a multiplicity of methods for the manufacture of medical devices that are thus appropriate, examples of which include, but are not limited to, foam processing, blow molding or film molding, sheet forming processes, profile extrusion, rotational molding, compression molding, thermoset pre-preg

15 processes, electrospinning or spraying, electrohydrodynamic spinning or spraying, and reaction injection molding processes.

Furthermore, medical devices in accordance with the present invention can be provided with enhanced properties or capabilities due to the inclusion of at least one nanomaterial therein, which properties and capabilities, in turn, are expected to

20 render the inventive medical devices particularly advantageous to use in therapeutic treatments. As a result, the present invention also provides methods of medical care utilizing the inventive devices. Generally speaking, the inventive method comprises providing a medical device comprising at least one nanomaterial, and utilizing the device to perform a diagnostic or therapeutic treatment. In those embodiments of

25 the invention wherein the device is a device intended to be introduced into a cardiovascular lumen, the method may further comprise inserting the device into the lumen and advancing it to the desired treatment site. As desired, the method may further comprise the delivery of the device to the treatment site, as may be the case with a stent, or performance of a desired treatment and removal of the device from

30 the lumen, as may be the case with an angioplasty catheter.

Advantageously, in embodiments where the systemic, controlled delivery of therapeutic agents is desired, the method may comprise the steps of delivering a

device followed either relatively immediately, or after a delay, by the systemic delivery of nanoparticles comprising a material capable of adhering to the delivered device, the therapeutic agent desirably delivered, and a means for causing the delivery of the therapeutic agent from the nanoparticles to the treatment site, e.g., a 5 biodegradable material.

The invention will now be further illustrated in the following examples, which are not intended to be limiting, but rather, have been chosen and described so that others skilled in the art may appreciate and understand the principles and practices of the present invention.

10

Example 1

Preparation of inner shaft catheter tubing with an HDPE/POSS nanocomposite

1) Preparation of the HDPE/POSS nanocomposite by twin-screw extrusion compounding

15 An organically functionalized POSS (MS0830, an OctaMethyl-POSS commercially available from Hybrid Plastics, Fountain Valley, CA) was compounded with high density polyethylene (HDPE Marlex 4903, commercially available from Chevron-Phillips Chemical Company, Houston, TX). In particular, a material feed ratio of HDPE to POSS of 4:1 was fed into a counter rotating 20 dispersive twin screw compounder ZSE 27 (commercially available from Leistritz Company, Allendale, NJ) operating at 190°C and a speed of 200 RPM. The compounding output was at 5 pounds per hour.

2) Extrusion of inner shaft catheter tubing incorporating HDPE/POSS 25 nanocomposite material.

A 4:1 mixture of the HDPE/POSS nanocomposite to plexar 390 anhydride modified polyethylene (commercially available from Equistar Chemical Company, Houston, TX) was premixed and further diluted at a 3:1 ratio with Marlex 4903 polyethylene and extruded into tubing of dimensions of 0.018 inch x 0.024 inch at 30 220°C.

Example 2**Preparation of outer shaft catheter tubing with a Pebax®/POSS nanocomposite****1) Preparation of the Pebax®/POSS nanocomposite by twin-screw extrusion compounding**

5 An organically functionalized POSS (AM0265, an Aminopropylisobutyl-POSS commercially available from Hybrid Plastics) was compounded with Pebax®7233 (Pebax® is a polyether block amide commercially available from Atofina, Brussels, Belgium). In particular, a material feed ratio of Pebax® to POSS of 4:1 was fed into a counter rotating dispersive Leistritz ZSE 27 twin screw
10 compounder operating at 200°C and a speed of 100 RPM. The compounding output was at 5 pounds per hour.

2) Extrusion of outer shaft catheter tubing incorporating Pebax®/POSS nanocomposite material

15 A 3:1 dilution of the Pebax®/POSS nanocomposite to Pebax®7233 was prepared and extruded into outer shaft tubing with dimensions of 0.0306 inch x 0.0362 inch at 226°C

It is expected that, during the tubing extrusion process, the nanocomposite will be more stable than conventional filled Pebax®. It is further believed that, if
20 the tubing produced by this method were subject to an EtO sterilization, that the POSS nanofiller will reduce or substantially prevent the oriented Pebax® chains from relaxing to a detrimental degree, as compared to such relaxation that would be expected to occur in an unfilled Pebax® medical device or device component when subjected to such sterilizing treatment.

25

Example 3**Preparation of outer shaft catheter tubing with a Pebax®/Clay nanocomposite**

A Pebax®/Clay nanocomposite material said to contain 95% Pebax® 7233 and 5% Clay filler with the trade designation of 2099 X 83109 C was purchased
30 from RTP Company (Winona, MN). The material was extruded into acceptable outer shaft tubing with dimensions 0.0306 inch x 0.0362 inch at an extrusion temperature of 226°C.

Example 4**Preparation of multilayer tubing with a Pebax®/montmorillonite nanocomposites**

5 A Pebax®/montmorillonite nanocomposite material containing 95% of a 72 durometer Pebax® material (such as Pebax® 7233 commercially available from Atochem) and 5% montmorillonite filler will be compounded with a twin screw extruder as described above. The nanocomposite material will then be coextruded with non-filled Pebax® at a temperature sufficient to provide appropriate viscosity 10 for extrusion, i.e., from about 190° C to about 215° C, into acceptable trilayer tubing having the Pebax®/montmorillonite nanocomposite as a middle layer and non-filled Pebax® as the inner and outer layers. The trilayer tubing will have dimensions appropriate for the intended use of the tubing. If the tubing is to be used, e.g., in the formation of a balloon, suitable dimensions would be an inner diameter of about 15 0.0176 inch and an outer diameter of about 0.342 inch.

Example 5**Preparation of monolayer tubing with a Pebax®/Modified montmorillonite nanocomposites**

20 A Pebax®/montmorillonite nanocomposite material containing 90% of a 70 durometer Pebax® material (such as Pebax® 7033 commercially available from Atochem) and 10% modified montmorillonite filler will be compounded with a twin screw extruder as described above. Prior to compounding, the montmorillonite will be modified with a functionalizer comprising a block copolymer capable of 25 interacting with polyether and/or polyamide, as described hereinabove. The nanocomposite material will be extruded at a temperature sufficient to provide appropriate viscosity for extrusion, i.e., from about 190° C to about 215° C, into acceptable monolayer tubing having dimensions appropriate for the intended use of the tubing. This tubing can then be used to form balloons, the inner lumen of catheters, the outer lumen of catheters, and the like. If the tubing is to be used, e.g., 30 in the formation of a balloon, suitable dimensions would be an inner diameter of about 0.0176 inch and an outer diameter of about 0.342 inch.

Example 6**Preparation of monolayer tubing with a Nylon 12/Modified montmorillonite nanocomposites**

5 A nylon 12/montmorillonite nanocomposite material containing 99% of a nylon 12 (commercially available under the tradename Rilsan® from Atofina) and 1% Modified montmorillonite filler will be prepared as follows. All materials will either be purchased as powders or ground into powders by any known method. The montmorillonite will be modified with a functionalizer comprising block polyamide 10 or any material having polyamide groups, as described hereinabove. The powdered nylon 12 and powdered functionalized montmorillonite will be mixed together and fed into an extrusion process via a gravimetric feeding device (or any other acceptable powder feeding mechanism). The nanocomposite material will then be extruded at a temperature sufficient to provide appropriate viscosity for extrusion, 15 i.e. from about 210° C to about 240° C, typically 220° C to 230° C, into acceptable monolayer tubing having dimensions appropriate for the intended use of the tubing. Such uses could include, e.g., formation of balloons, inner lumens of catheters, outer lumens of catheters, etc. Tubing comprising such a nanocomposite is contemplated to be particularly useful in the formation of balloons, for which use appropriate 20 tubing dimensions are an inner diameter of about 0.0176 inch and an outer diameter of about 0.342 inch.

Example 7**Preparation of monolayer tubing with a polyurethane/carbon nanofiber nanocomposites**

25 Polyacrylonitrile will be electrospun about a rod or tube of the desired tubing dimensions. The resulting polyacrylonitrile fibers will be heated in air, causing it to oxidize. The resulting modified polyacrylonitrile fibers will then be heated in the absence of air to a temperature greater than at least about 1300°C. This process will causes the carbonization of the fibers into the form of ribbon like fused ring 30 polymer, which will then condenses to provide a lamellar basal plane structure of nearly pure carbon.

A polyurethane will be dissolved in an amount of tetrahydrofuran. The carbon nanofiber structure will be dipcoated with the polyurethane/tetrahydrofuran solution multiple times so as to create a polyurethane matrix surrounding the carbon nanofiber structure, thereby forming acceptable monolayer tubing having 5 dimensions appropriate for the intended use of the tubing. This tubing can then be used to form balloons, the inner lumen of catheters, the outer lumen of catheters, and the like. If the tubing is to be used, e.g., in the formation of a balloon, suitable dimensions would be an inner diameter of about 0.0176 inch and an outer diameter of about 0.342 inch.

10

Example 8

Preparation of an embolic filter with a Pellethane®/functionalized clay nanocomposite

A suitable Pellethane® will be dissolved in a corresponding suitable amount of tetrahydrofuran. An amount of alkylammonium-modified montmorillonite, up to 15 40%, will be added to the Pellethane® solution, and the solution mixed in a high speed mixer operatively disposed relative to an ultrasonic transducer until microscopic inspection reveals that the clay is dispersed and that no clots thereof remain.

The Pellethane®/montmorillonite nanocomposite will then be 20 electrostatically sprayed onto a suitable mold until the desired thickness is achieved. In particular, it is believed that an embolic filter so produced and having a thickness of about 0.0003", about 4 times less thick than a filter comprising a traditional material, will yet exhibit up to a 300% increase in mechanical strength over filters prepared from conventional materials.

25

Other embodiments of this invention will be apparent to those skilled in the art upon consideration of this specification or from practice of the invention disclosed herein. Various omissions, modifications, and changes to the principles and embodiments described herein may be made by one skilled in the art without departing from the true scope and spirit of the invention which is indicated by the 30 following claims.

What is claimed is:

- 1) A medical device comprising at least one nanomaterial.
- 5 2) The medical device of claim 1, wherein the nanomaterial comprises at least one plurality of nanoparticles.
- 3) The medical device of claim 2, wherein the nanoparticles comprise carbon nanotubes or nanofibers.
- 10 4) The medical device of claim 3, wherein the medical device comprises multiple layers and the carbon nanotubes or nanofibers are interspersed between two adjacent layers.
- 15 5) The medical device of claim 4, wherein the medical device is catheter shafting.
- 6) The medical device of claim 1, wherein the nanomaterial comprises a nanocomposite material further comprising at least one plurality of nanoparticles and a matrix material.
- 20 7) The medical device of claim 6, wherein the device is an intralumen cardiovascular medical device.
- 25 8) The medical device of claim 7, wherein the matrix material comprises a thermoplastic material and the nanoparticles comprise a clay.
- 9) The medical device of claim 8, wherein the matrix material comprises a polyurethane and the clay comprises montmorillonite.
- 30 10) The medical device of claim 9, wherein the montmorillonite is functionalized with alkylammonium.

- 11) The medical device of claim 8, wherein the medical device comprises an embolic filter.
- 5 12) A medical care kit, comprising at least one medical device and at least one nanomaterial provided in packaging.
- 13) The medical care kit of claim 12, wherein the at least one nanomaterial is included in at least a component of the at least one medical device.
- 10 14) The medical care kit of claim 12, wherein the nanomaterial comprises a first plurality of nanoparticles.
- 15 15) The medical care kit of claim 14, wherein the first plurality of nanoparticles comprise at least one biodegradable, biocompatible material, and at least one therapeutic agent.
- 16) The medical care kit of claim 15, wherein the first plurality of nanoparticles further comprise a magnetic material.
- 20 17) The medical care kit of claim 15, wherein the first plurality of nanoparticles further comprise a heat sensitive material.
- 25 18) The medical care kit of claim 13, wherein the medical device is a stent, the at least one nanomaterial comprises a coating applied to at least a portion of the stent and wherein the nanomaterial coating comprises a thermoplastic matrix material and magnetic nanoparticles.
- 30 19) A method of medical care comprising bringing a medical device into therapeutic contact with a body to effectuate a treatment or diagnosis thereof, wherein the medical device comprises at least one nanomaterial.

20) The method of claim 19, wherein the method of medical care is a method of treatment.

21) The method of claim 20, wherein the medical device is a transluminal 5 medical device.

22) The method of claim 21, wherein the medical device is a stent delivery catheter.

10 23) The method of claim 22, wherein the stent delivery catheter is caused to deliver a stent to a desired treatment site.

24) The method of claim 23, further comprising causing a plurality of nanoparticles to be delivered to the treatment site.

15 25) The method of claim 24, wherein the nanoparticles operatively disposed relative to the stent at the time of delivery of the stent.

26) The method of claim 25, wherein the nanoparticles are a component of a nanomaterial coating provided on at least a portion of a surface of the stent.

20 27) The method of claim 24, wherein the nanoparticles are caused to be operatively disposed relative to the desired treatment site at substantially the same time as the stent is delivered.

25 28) The method of claim 24, wherein the nanoparticles are caused to be operatively disposed relative to the desired treatment site after the stent has been delivered.

30 29) The method of claim 28, wherein the step of causing the nanoparticles to be operatively disposed relative to the desired treatment site is repeated a plurality of times.

30) A method of making at least a component of a medical device, comprising:
providing a nanomaterial;
preparing at least a component of the medical device with the nanomaterial.

5 31) The method of claim 30, wherein the nanomaterial comprises a first plurality
of nanoparticles.

32) The method of claim 31, wherein the nanoparticles comprise carbon
nanotubes, nanofibers, or nanowires.

10 33) The method of claim 32, wherein the nanomaterial is provided by
electrospinning or electrohydrodynamically spraying a material onto a mold.

15 34) The method of claim 33, wherein the material is a carbonizable material that
is electrohydrodynamically spun unto a mandrel and then carbonized to provide
carbon nanotubes, nanowires, or nanofibers.

20 35) The method of claim 34, wherein the component of the medical device is
provided by depositing a matrix material onto the mandrel and about at least a
portion of the carbon nanoparticles.

36) The method of claim 32, wherein the component of the device prepared is a
tie layer provided between a first and a second layer of the medical device.

25 37) The method of claim 36, wherein the tie layer is provided by causing at least a
portion of the first layer of the medical device to come into contact with a solution
comprising the carbon nanoparticles.

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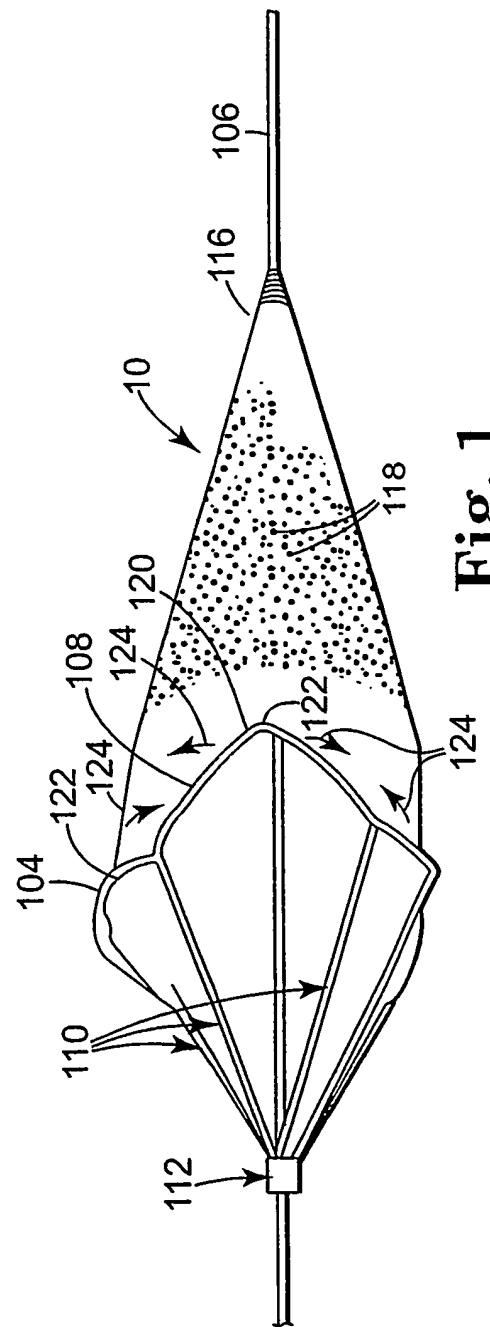


Fig. 1 -

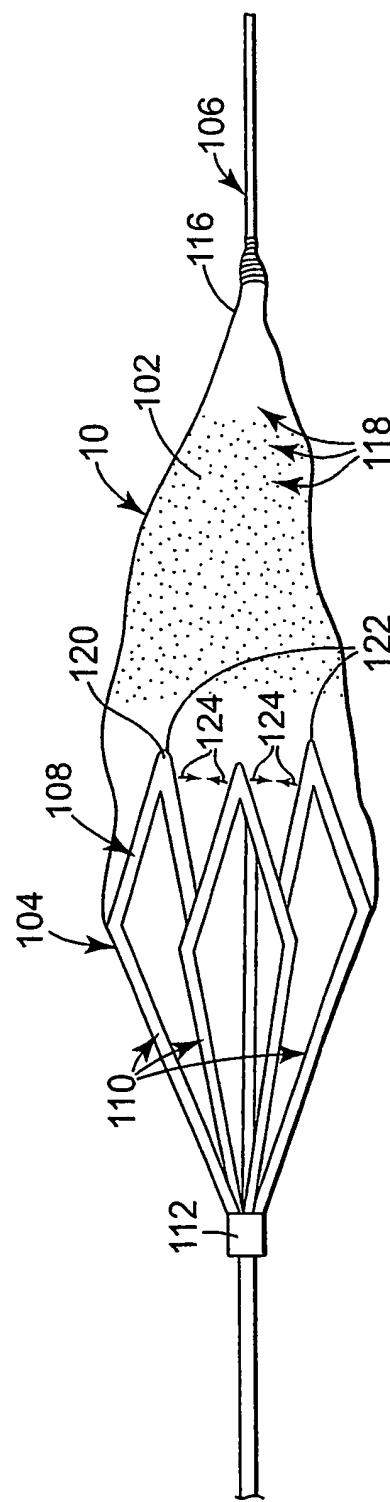


Fig. 2

